2-21-06 07/044 086 C-of-C.

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

PATENT NO.	:	4,959,366
APPLICATION NO.	:	44,086
ISSUE DATE	:	September 25, 1990
INVENTORS	:	Cross, et al.

RESUBMISSION OF REQUEST FOR CERTIFICATE OF CORRECTION

ATTN: Certificate of Correction Branch
Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Sir:

Enclosed is a Request for Certificate of Correction (including the Certificate and Exhibits) for U.S. Patent No. 4,959,366. Applicants previously filed this Request for Certificate of Correction with the Office and are re-submitting the Request for Certificate of Correction at the instruction of the Office. The relevant facts are as follows:

(1) On April 27, 2005, the Office issued a Certificate Extending Patent Term Under 35 U.S.C. § 156 for U.S. Patent No. 4,959,366 (the "Certificate Extending Patent Term").

(2) The Certificate Extending Patent Term contained several errors as outlined in the attached Request for Certificate of Correction.

(3) In May, 2005, the undersigned attorney contacted Karin Ferriter of the Patent Extension Branch regarding the errors in the Certificate Extending Patent Term. Ms. Ferriter instructed the undersigned to send the Request for Certificate of Correction to her by facsimile for filing, and indicated that she would forward the Request, along with the file to the Certificate of Correction Branch.

(4) On June 1, 2005, the attached Request for Certificate of Correction was sent to Ms. Ferriter by facsimile. A copy of the facsimile confirmation report is attached.

(5) In January, 2006, the undersigned attorney contacted the Certificate of Correction Branch to determine the status of the Request for Certificate of Correction and was advised that the Request for Certificate of Correction had not been received by the Certificate of Correction Branch. The individual with whom the undersigned spoke advised Applicant to resubmit the Request for Certificate of Correction to the Certificate of Correction Branch to ensure proper consideration thereof.

Accordingly, applicants are now re-submitting the previously filed Request for Certificate of Correction (including the Certificate and Exhibits) for U.S. Patent No. 4,959,366, and respectfully request issuance of a Certificate of Correction.

If the Office has any questions please contact the undersigned at the telephone number indicated.

Respectfully submitted,

Daniel S. Kasten
Registration No. 45,363
Telephone: 314-274-5218

Pharmacia Corporation of Pfizer Inc.
P.O. Box 1027
St. Louis, MO 63006

****
*** TX REPORT ***
****

TRANSMISSION OK

TX/RX NO	3623
CONNECTION TEL	915712737744
SUBADDRESS	
CONNECTION ID	
ST. TIME	06/01 14:33
USAGE T	06'28
PGS. SENT	15
RESULT	OK

CERTIFICATE OF TRANSMISSION BY FACSIMILE (37 CFR 1.8) Applicant(s): Cross et al.			Docket No. PCL428
Application No. 44,086 Patent# 4,959,336	Filing Date September 25, 1990	Examiner n/a	Group Art Unit n/a

Invention: ANTI-ARRHYTHMIC AGENTS

I hereby certify that this **Certificate of Correction; Request for Certificate of Correction; Exhibits A-H**
(Identify type of correspondence)
is being facsimile transmitted to the United States Patent and Trademark Office (Fax. No. **571.273.7744**
on **June 1, 2005**
(Date)

Linda K. Cooper
(Typed or Printed Name of Person Signing Certificate)

Linda K. Cooper
(Signature)

15 pages exc

Note: Each paper must have its own certificate of mailing.

FEB 06 2006

CERTIFICATE OF TRANSMISSION BY FACSIMILE (37 CFR 1.8) Applicant(s): Cross et al.			Docket No. PCL428
Application No. 44,086 Patent# 4,959,336	Filing Date September 25, 1990	Examiner n/a	Group Art Unit n/a

Invention: ANTI-ARRHYTHMIC AGENTS

I hereby certify that this Certificate of Correction; Request for Certificate of Correction; Exhibits A-H
(Identify type of correspondence)
is being facsimile transmitted to the United States Patent and Trademark Office (Fax. No. 571.273.7744

on June 1, 2005
(Date)

Linda K. Cooper
(Typed or Printed Name of Person Signing Certificate)

Linda K. Cooper
(Signature)

15 pages lkc

Note: Each paper must have its own certificate of mailing.

PTO/SB/44 (04-04)
Approved for use through 04/30/2007. OMB 0651-0033
U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO : **4,959,366**

DATED : **September 25, 1990**

INVENTOR(S) : **Cross et al.**

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

In the Certificate Extending Patent Term Under 35 U.S.C. § 156, issued April 27, 2005:

(a) in the heading of the Certificate, following [(45) ISSUED], "May 25, 1990" should read --September 25, 1990--.

(b) in the heading of the Certificate, following [(95) PRODUCT], "TIKOSYN® (methanesulfonamide)" should read -- TIKOSYN® (dofetilide)--.

(c) at lines 3-4 of the first paragraph of the text of the Certificate, "TIKOSYN® (methanesulfonamide)" should read -- TIKOSYN® (dofetilide)--.

MAILING ADDRESS OF SENDER: **Daniel S. Kasten**
P.O. Box 1027
Chesterfield, Missouri 63006

PATENT NO. **4,959,336**

No. of additional copies

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

PATENT NO.	:	4,959,366
APPLICATION NO.	:	44,086
ISSUE DATE	:	September 25, 1990
INVENTORS	:	Cross, et al.

REQUEST FOR CERTIFICATE OF CORRECTION

ATTN: Certificate of Correction Branch
Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Sir:

Applicant requests the Office to issue a Certificate of Correction pursuant to 37 C.F.R. 1.322 for U.S. Patent 4,959,366. Specifically, Applicant requests the Office to correct the Certificate Extending Patent Term Under 35 U.S.C. § 156 issued April 27, 2005, for U.S. Patent 4,959,366. The requested corrections are identified in the attached Certificate of Correction Form PTO/SB/44.

Applicant believes that the errors to be corrected are attributable solely to the Office and requests expedited processing and grant of the Certificate of Correction. If the Office disagrees and believes that one or more of the errors are attributable to Applicant, the Office may instead treat this request as one for the Office to issue a Certificate of Correction pursuant to 37 C.F.R. 1.323 for U.S. Patent 4,959,366.

The specific facts related to the requested corrections are set forth below:

(1) The heading of the Certificate Extending Patent Term lists the incorrect issue date for U.S. Patent 4,959,366. The correct issue date is clearly reflected on the face of U.S. Patent 4,959,366. Applicant is not aware of any prior occurrences of this error. Thus, it appears that the error is of a

clerical nature, and attributable solely to the Office. A copy of the face sheet of the '366 patent is attached as Exhibit A.

(2) The remaining two errors in the Certificate Extending Patent Term are references to the product as "TIKOSYN® (methanesulfonamide)" rather than as "TIKOSYN® (dofetilide)".

(3) The active ingredient in TIKOSYN® is *N*-[4-[2-[methyl[2-[4-[(methylsulfonyl)amino]phenoxy]ethyl]amino]ethyl]phenyl]-methanesulfonamide. Dofetilide is the U.S. Adopted Name and the International Nonproprietary Name for this compound. On November 18, 1999, Applicant submitted an application for extension of the term of U.S. Patent 4,959,366 under 35 U.S.C. § 156. The application for extension referred to the approved product "TIKOSYN® (dofetilide)". A copy of the first page of the application is attached as Exhibit B.

(4) By letter to the Food and Drug Administration ("FDA") on or about December 1, 1999, the Office requested FDA to confirm "that the product identified in the application, TIKOSYN® (dofetilide), has been subject to a regulatory review period within the meaning of 35 U.S.C. § 156(g) before its first commercial marketing or use and that the application for patent term extension was filed within the sixty-day period after the product was approved." A copy of this letter is attached as Exhibit C.

(5) By letter to the Office on or about April 26, 2000, the FDA confirmed that the product had been subject to a regulatory review period and that the subsequent marketing and use represented the first commercial marketing or use of that product. The FDA further confirmed that the application for patent term extension was timely filed. Although this letter listed the correct product trade name (TIKOSYN®), New Drug Application (NDA No. 20-931) and patent (U.S. Patent 4,959,366), the generic name listed for the product was incorrect (methanesulfonamide rather than dofetilide). A copy of this letter is attached as Exhibit D.

(6) By letter to the Office on or about November 17, 2003, the FDA indicated that it had published its determination of the product's regulatory review

period and that it considered the regulatory review period determination to be final. This letter listed the correct product trade name (TIKOSYN®), New Drug Application (NDA No. 20-931) and patent (U.S. Patent 4,959,366), but the generic name listed for the product was incorrect (methanesulfonamide rather than dofetilide). A copy of this letter is attached as Exhibit E.

(7) On January 25, 2005, the Office issued a Notice of Final Determination stating that U.S. Patent No. 4,959,366 was entitled to a five-year extension of term through September 25, 2012. Although the Notice of Final Determination listed the correct patent (U.S. Patent 4,959,366) and product trade name (TIKOSYN®), the generic name for the product was listed incorrectly three times (methanesulfonamide rather than dofetilide). A copy of the Notice of Final Determination is attached as Exhibit F.

(8) On January 26, 2005, Applicant filed a letter stating that Pfizer Inc, the owner of record of U.S. Patent No. 4,959,366, did not wish to file a request for reconsideration of the final determination as to the length of extension of the term of that patent. This letter did not expressly comment on the incorrect references to the generic name (methanesulfonamide rather than dofetilide) contained in the Notice of Final Determination. A copy of this letter is attached as Exhibit G.

(9) On April 27, 2005, the Office issued a Certificate Extending Patent Term Under 35 U.S.C. § 156 for U.S. Patent 4,959,366. Although the Certificate Extending Patent Term listed the correct patent (U.S. Patent 4,959,366) and product trade name (TIKOSYN®), the generic name for the product was listed incorrectly two times (methanesulfonamide rather than dofetilide). A copy of the Certificate Extending Patent Term is attached as Exhibit H.

Accordingly, Applicant respectfully requests expedited handling and grant of a Certificate of Correction pursuant to 37 C.F.R. 1.322 for U.S. Patent 4,959,366. In the event the Office believes, however, that one or more of the

errors to be corrected are attributable to Applicant, Applicant respectfully requests the Office instead to treat this request as one for issuance of a Certificate of Correction pursuant to 37 C.F.R. 1.323 for U.S. Patent 4,959,366. The Office is hereby authorized to charge any fees that may be required in this matter to Deposit Account No. 19-1025.

Respectfully submitted,

Daniel S. Kasten
Registration No. 45,363
Telephone: 314-274-5218

Pharmacia Corporation of Pfizer Inc.
P.O. Box 1027
Chesterfield, Missouri 63006

United States Patent [19]

Cross et al.

[11] Patent Number: 4,959,366

[45] Date of Patent: Sep. 25, 1990

[54] ANTI-ARRHYTHMIC AGENTS

[75] Inventors: Peter E.Cross, Geoffrey N. Thomas, John E. Arrowsmith, all of New York, N.Y.

[73] Assignee: Pfizer Inc., New York, N.Y.

[21] Appl. No.: 44,086

[22] Filed: Apr. 29, 1987

[30] Foreign Application Priority Data

May 1, 1986 [GB] United Kingdom 8610668
Dec. 17, 1986 [IE] Ireland 8630059

[51] Int. Cl.[3] C07C 143/75; C07C 93/06

[52] U.S. Cl. 514/239.5; 514/255; 514/315; 514/410; 514/605; 514/618; 514/619; 514/620; 514/621; 514/649; 514/651; 514/654; 544/165; 544/167; 544/399; 546/226; 546/539; 564/99; 564/162; 564/164; 564/166; 564/336; 564/341; 564/348; 564/354

[58] Field of Search 564/99, 336, 341, 348, 564/354, 162, 164, 166; 514/605, 821, 239.5, 255, 315, 410, 618, 619, 620, 621; 546/226; 548/539; 544/399, 165, 167

[56] References Cited

U.S. PATENT DOCUMENTS

3,341,584	9/1967	Larsen et al.	260/556
3,478,149	11/1969	Larsen et al.	424/228
3,574,741	4/1971	Gould et al.	260/556
3,660,487	5/1972	Larsen et al.	260/556
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0164865	12/1985	European Pat. Off.	514/605
977261	7/1966	France	514/605
1263987	2/1972	United Kingdom	514/605
1301134	12/1982	United Kingdom	514/605
2135883	9/1984	United Kingdom	514/605

OTHER PUBLICATIONS

Larsen et al., "Sulfonanilides, II Anbalogs of Catecholamines", J. Am. Chem. Soc. 75, 4334 (1953).

Primary Examiner—Richard L. Raymond
Assistant Examiner—Raymond Covington
Attorney, Agent, or Firm—Peter C. Richardson; Paul H. Ginsburg; Robert F. Sheyka

[57] ABSTRACT

A series of [N-alkyl-N-(nitro-, alkylsulphonamido, or amino-phenalkyl)amino]-alkyl, alkoxy or alkylthio phenyl derivatives having utility as anti-arrhythmic agents.

48 Claims, No Drawings

Exhibit A

PC7068AADO

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

IN RE:	**U.S. Patent No. 4,959,366:**
ISSUED:	**September 25, 1990** :
TO:	**Peter E. Cross, et al.** :
FOR:	**Anti-Arrhythmic Agents** :
FROM:	**Serial No. 44,086** :
FILED:	**APRIL 29, 1987** :

BOX PATENT EXTENSION
Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

Your applicant, Pfizer Inc. ("PFIZER"), a corporation of the State of Delaware and having a place of business at 235 East 42nd Street, New York, New York, represents that it is the legal owner of record of Letters Patent of the United States No. 4,959,366, granted to Peter E. Cross, Geoffrey N. Thomas and John E. Arrowsmith[1] on the twenty-fifth day of September, 1990, for Anti-Arrhythmic Agents, by virtue of an assignment, recorded in the United States Patent and Trademark Office on the twenty-ninth day of April, 1987, at Reel 4780, Frame 0801; that Pfizer Pharmaceuticals Production Corporation Limited ("PPPCL") is an indirectly-owned subsidiary of PFIZER; that PFIZER controls, indirectly, all the voting shares of PPPCL; that PPPCL is the owner of New Drug Application ("NDA") No. 20-931 for TIKOSYN (dofetilide), claimed by U.S. Patent No. 4,959,366; and that PFIZER is entitled to rely on the marketing approval for TIKOSYN (dofetilide) arising from NDA No. 20-931.

[1] Subsequently, a certificate of correction (Exhibit B) dated May 11, 1999 certified that the correct inventorship is: Peter E. Cross and Geoffrey N. Thomas.

DEC - 1 1999

UNITED S ES DEPARTMENT OF COMMERCE
Patent and Trademark Office
ASSISTANT SECRETARY AND COMMISSIONER
OF PATENTS AND TRADEMARKS
Washington, D.C. 20231

2C 7063 RFS

David T. Read
Acting Director Regulatory Policy Staff, CDER
Food and Drug Administration
1451 Rockville Pike, HFD-7
Rockville, MD 20852

Dear Mr. Read:

The attached application for patent term extension of U.S. Patent No. 4,959,366 was filed on November 19, 1999, under 35 U.S.C. § 156.

The assistance of your Office is requested in confirming that the product identified in the application, TIKOSYN (dofetilide), has been subject to a regulatory review period within the meaning of 35 U.S.C. § 156(g) before its first commercial marketing or use and that the application for patent term extension was filed within the sixty-day period after the product was approved. Since a determination has not been made whether the patent in question claims a product, or the method of use of manufacturing such a product, which has been subject to the Federal Food, Drug and Cosmetic Act, this communication is NOT to be considered as notice which may be made in the future pursuant to 35 U.S.C. § 156(d)(2)(A).

Our review of the application to date indicates that the subject patent would be eligible for extension of the patent term under 35 U.S.C. § 156.

Inquiries regarding this communication should be directed to the undersigned at (703) 306-3159 (telephone) or (703)308-6916 (facsimile).

Karin Tyson
Legal Advisor
Special Program Law Office
Office of the Deputy Assistant Commissioner
for Patent Policy and Projects

cc: Gregg C. Benson
Pfizer Inc.
Patent Department
Eastern Point Road
Groton CT 06340

kt

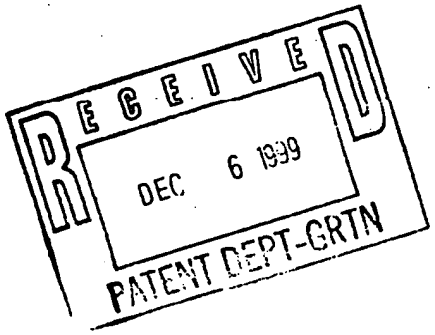


Exhibit C

DEPARTMENT OF HEALTH & HUMAN SERVICES — Public Health Service

Food and Drug Administration
Rockville MD 20857

APR 26 2000

Re: Tikosyn®
Docket No. 00N-1248

The Honorable Q. Todd Dickinson
Director of U.S. Patent and Trademark Office
Commissioner for Patents
Box Pat. Ext.
Washington, D.C. 20231

Dear Director Dickinson:

This is in regard to the application for patent term extension for U.S. Patent No. 4,959,366 filed by Pfizer under 35 U.S.C. § 156. The human drug product claimed by the patent is Tikosyn® (methanesulfonamide), which was assigned new drug application (NDA) No. 20-931.

A review of the Food and Drug Administration's official records indicates that this product was subject to a regulatory review period before its commercial marketing or use, as required under 35 U.S.C. § 156(a)(4). Our records also indicate that it represents the first permitted commercial marketing or use of the product, as defined under 35 U.S.C. § 156(f)(1), and interpreted by the courts in *Glaxo Operations UK Ltd. v. Quigg*, 706 F. Supp. 1224 (E.D. Va. 1989), *aff'd*, 894 F. 2d 392 (Fed. Cir. 1990).

The NDA was approved on October 1, 1999, which makes the submission of the patent term extension application on November 19, 1999, timely within the meaning of 35 U.S.C. § 156(d)(1).

Should you conclude that the subject patent is eligible for patent term extension, please advise us accordingly. As required by 35 U.S.C. § 156(d)(2)(A) we will then determine the applicable regulatory review period, publish the determination in the *Federal Register*, and notify you of our determination.

Please let me know if we can be of further assistance.

Sincerely yours,

Jane A. Axelrad
Associate Director for Policy
Center for Drug Evaluation and Research

cc: Gregg C. Benson
Pfizer, Inc.
Patent Department
Eastern Point Rd
Groton, CT 06340

Exhibit D

PC007008A RAP

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Former # PLC 428

Food and Drug Administration
Rockville MD 20857

Re: Tikosyn
Docket No. 00N-1248

The Honorable James E. Rogan
Under Secretary of Commerce for Intellectual Property and
Director of the United States Patent and Trademark Office
Box Pat. Ext.
P.O. Box 1450
Alexandria, VA 22313-1450

NOV 17 2003

Dear Director Rogan:

This is in regard to the patent term extension application for U.S. Patent No. 4,959,366 filed by Pfizer under 35 U.S.C. § 156. The patent claims Tikosyn (methanesulfonamide), NDA 20-931.

In the April 18, 2003, issue of the Federal Register (68 Fed. Reg. 19212), the Food and Drug Administration published its determination of this product's regulatory review period, as required under 35 U.S.C. § 156(d)(2)(A). The notice provided that on or before October 15, 2003, 180 days after the publication of the determination, any interested person could file a petition with FDA under 35 U.S.C. § 156(d)(2)(B)(i) for a determination of whether the patent term extension applicant acted with due diligence during the regulatory review period.

The 180-day period for filing a due diligence petition pursuant to this notice has expired and FDA has received no such petition. Therefore, FDA considers the regulatory review period determination to be final.

Please let me know if we can provide further assistance.

Sincerely yours,

Jane A. Axelrad

Jane A. Axelrad
Associate Director for Policy
Center for Drug Evaluation and Research

cc: Gregg C. Benson
Pfizer, Inc.
Patent Department
Eastern Point Rd.
Groton, CT 06340

RECEIVED

JAN 2 1 2004

PHARMACIA
GLOBAL PATENT DEPT.
ST. LOUIS, MO

Exhibit E

UNITED STATES PATENT AND TRADEMARK OFFICE

Commissioner for Patents
United States Patent and Trademark Office
P.O. Box 1450
Alexandria, VA 22313-1450
www.uspto.gov

Mailed: January 25, 2005

Gregg C. Benson
Pfizer, Inc.
Patent Department
Eastern Point Road
Groton CT 06340

Re: Patent Term Extension
Application for
U.S. Patent No. 4,959,366

NOTICE OF FINAL DETERMINATION

A determination has been made that U.S. Patent No. 4,959,366, which claims the human drug product TIKOSYN® (methanesulfonamide) and a method of use of said product, is eligible for patent term extension under 35 U.S.C. § 156. The period of extension has been determined to be five years.

A single request for reconsideration of this final determination as to the length of extension of the term of the patent may be made if filed within one month of the date of this notice. Extensions of time under 37 CFR § 1.136(a) are not applicable to this time period. In the absence of such request for reconsideration, the Director will issue a certificate of extension, under seal, for a period of five years.

Under 35 U.S.C. § 156(c), the period of extension is calculated using the Food and Drug Administration's determination of the length of the regulatory review period as published in the Federal Register of April 18, 2003 (68 Fed. Reg. 19212).

Period of Extension	=	½ (Testing Phase) + Approval Phase
	=	½ (2,778 - 56) + 572
	=	1,933 days

Since the regulatory review period began August 1, 1990, before the patent issue date (September 25, 1990), the period during the regulatory review period on or before the issue date of the patent has been subtracted from the "testing phase" in the above determination. No determination of a lack of due diligence under 35 U.S.C. § 156(c)(1) was made.

The five year limitation of 35 U.S.C. § 156(g)(6)(A) applies in the present situation, however, because the patent was issued after the date of enactment (September 24, 1984) of 35 U.S.C. § 156. Since the period of extension calculated under 35 U.S.C. § 156(c) cannot exceed five years under 35 U.S.C. § 156(g)(6)(A), the period of extension will be for five years.

The 14 year limitation of 35 U.S.C. § 156(c)(3) does not operate to further reduce the period of extension determined above.

Upon issuance of the certificate of extension, the following information will be published in the Official Gazette:

U.S. Patent No.:	4,959,366
Granted:	September 25, 1990
Original Expiration Date[1]:	September 25, 2007
Applicant:	Cross, et al.

[1]Subject to the provisions of 35 U.S.C. § 41(b).

Owner of Record:	Pfizer Inc.
Title:	ANTI-ARRHYTHMIC AGENTS
Classification:	514/239.5
Product Trade Name:	TIKOSYN® (methanesulfonamide)
Term Extended:	Five years
Expiration Date of Extension:	September 25, 2012

Any correspondence with respect to this matter should be addressed as follows:

By mail: Commissioner for Patents
Mail Stop Patent Ext.
P.O. Box 1450
Alexandria, VA 22313-1450

By FAX: (703) 273-7744
Attn: Office of Patent Legal Administration

Telephone inquiries related to this determination should be directed to the undersigned at 571-272-7744.

Karin Ferriter
Senior Legal Advisor
Office of Patent Legal Administration
Office of the Deputy Assistant Commissioner
for Patent Policy and Projects

cc: Office of Regulatory Policy
HFD - 13
5600 Fishers Lane
Rockville, MD 20857

Attention: Claudia Grillo

RE: TIKOSYN® (methanesulfonamide)
FDA Docket No.: 00N-1248

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

IN RE:	U.S. Patent No. 4,959,366	:
ISSUED:	September 25, 1990	:
TO:	Peter E. Cross, et al.	:
FOR:	Anti-Arrhythmic Agents	:
FROM:	Application Ser. No. 44,086	:
FILED:	April 29, 1987	

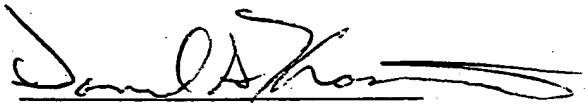
Commissioner for Patents
Mail Stop Patent Extension
P.O. Box 1450
Alexandria, VA 22313-1450

LETTER RE: CERTIFICATE OF EXTENSION

Sir:

Further to the Notice of Final Determination, dated January 25, 2005, Pfizer Inc., the Owner of Record of U.S. Patent No. 4,959,366, does NOT wish to file a request for reconsideration of the final determination as to the length of extension of the term of that patent. Accordingly, Pfizer Inc. respectfully requests that the Certificate of Extension be issued as soon as possible.

Respectfully submitted,



Daniel S. Kasten
Patent Counsel
Reg. No. 45,363
Tel: 314-274-5218

Pfizer Inc.
Patent Department
575 Maryville Centre Drive
St. Louis, MO 63141

Exhibit G

UNITED STATES PATENT AND TRADEMARK OFFICE

(12) CERTIFICATE EXTENDING PATENT TERM
UNDER 35 U.S.C. § 156

(68)	PATENT NO.	:	4,959,366
(45)	ISSUED	:	May 25, 1990
(75)	INVENTOR	:	Cross, et al.
(73)	PATENT OWNER	:	Pfizer Inc.
(95)	PRODUCT	:	TIKOSYN® (methanesulfonamide)

This is to certify that an application under 35 U.S.C. § 156 has been filed in the United States Patent and Trademark Office, requesting extension of the term of U.S. Patent No. 4,959,366 based upon the regulatory review of the product TIKOSYN® (methanesulfonamide) by the Food and Drug Administration. Since it appears that the requirements of the law have been met, this certificate extends the term of the patent for the period of

(94) Five Years

from September 25, 2007, the original expiration date of the patent, subject to the payment of maintenance fees as provided by law, with all rights pertaining thereto as provided by 35 U.S.C. § 156(b).

I have caused the seal of the United States Patent and Trademark Office to be affixed this 27th day of April 2005.

Jon W. Dudas
Under Secretary of Commerce for Intellectual Property and
Director of the United States Patent and Trademark Office

Exhibit H

CERTIFICATE OF MAILING BY "EXPRESS MAIL" (37 CFR 1.10) Applicant(s): **Cross et al.**				Docket No. **PCL428**
Application No. **44,086Patent4,959,336**	Filing Date **September 25,1990**	Examiner **n/a**	Customer No. **26648**	Group Art Unit **n/a**

Invention: **Anti-Arrhythmic Agents**

I hereby certify that the following correspondence:

Resubmission of Request for Cert. of Correction, Facsimile Transmission Receipt, Certificate of Transmission by Facsimile, Certificate of Correction, Request for Certificate of Correction (4pgs), Exhibit A,Exhibit B,Exhibit C, Exhibit D,Exhibit E, Exhibit F, Exhibit G, Exhibit H and return receipt postcard

(Identify type of correspondence)

is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 CFR 1.10 in an envelope addressed to: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450 on

February 17, 2006
(Date)

Connie Payne
(Typed or Printed Name of Person Mailing Correspondence)

Connie Payne
(Signature of Person Mailing Correspondence)

EV846966026 US
("Express Mail" Mailing Label Number)

Note: Each paper must have its own certificate of mailing.